Supplemental data

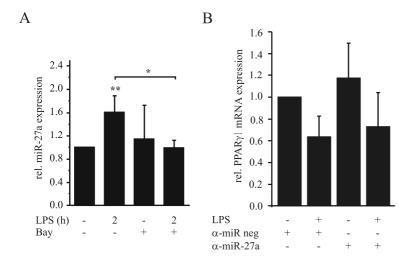
FIGURE LEGENDS

Suppl. Fig. S1. Modulation of miR-27a expression. A. MiR-27a expression was measured in primary human macrophages in response to LPS (1 μ g/ml, 2 h) by qPCR. To investigate a role of NFκB, cells were pre-stimulated for 1 h with 10 μ M Bay11-7082. Basal expression was set to 1. *B*. Primary human macrophages were transfected with 150 pmol of anti-miR-27a or negative control. After transfection, cells were stimulated with 1 μ g/ml LPS for 3 h and PPARγ1 mRNA level was determined by qPCR. Data present mean values \pm SE. n \geq 3. Statistics were analyzed with the paired Student's t-test. *p < 0.05, **p < 0.01.

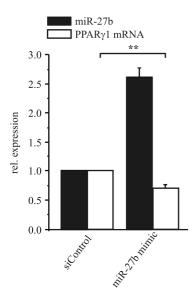
<u>Suppl. Fig. S2.</u> Effect of an approximately 2-fold increase of miR-27b on PPAR γ 1 expression. Primary human macrophages were transfected with miR-27b mimic, PPAR γ 1 mRNA and miR-27b expression was determined by qPCR. Data are given as mean values \pm SE, n = 3. Statistics were analyzed with the paired Student's t-test. **p < 0.01.

Suppl. Fig. S3. Impact of TNFα and IFNγ on PPARγ1 expression. *A*. Primary human macrophages were stimulated with 15 ng/ml TNFα or 10 U/ml IFNγ with or without LPS (1 µg/ml) for 3 h and PPARγ1 mRNA was measured by qPCR. *B*. miR-27b expression was measured by qPCR after stimulation of primary macrophages with 15 ng/ml TNFα for 2 h. *C*. Primary human macrophages were transfected with 150 pmol anti-miR-27b inhibitor and stimulated with 15 ng/ml TNFα for 3 h. PPARγ1 mRNA was determined by qPCR. Data present mean values \pm SE, n \geq 3. Statistics were analyzed using the paired Student's t-test.*p < 0.05, **p < 0.01, ***p < 0.001.

Suppl. Fig. S1



Suppl. Fig. S2.



Suppl. Fig. S3.

